

PROCESS FOR THE PREPARATION OF A METAL-ORGANIC COMPOUND
COMPRISING AT LEAST ONE IMINE LIGAND

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The invention relates to a process for the preparation of a metal-organic compound comprising at least one imine ligand according to formula 1. Metal-organic compounds thus produced are typically used as precatalyst in the production of polyolefins. Imine ligands for these precatalyst can be guanidine, iminoimidazoline, ketimine or phosphinimine, the manufacturing of which is described in WO 2070569, US 6114481 and US 6063879 respectively.

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The known production processes for phosphinimine comprising metal-organic compounds require at least two steps: (i) the synthesis of a N-trialkylsilyl substituted imine ligand, followed by (ii) contacting this ligand with a metal-organic precursor. However, in the one step process for the manufacturing of the imine ligand, as described in Z. Naturforschung. 29b, 328(1974) (the Staudinger reaction), azide chemistry is required. In this process, the most frequently used azide is azidotrimethylsilane, which is highly toxic and readily hydrolysable, releasing the highly toxic and both temperature and shock sensitive hydrazoic acid. Therefore, mixtures containing (partially) hydrolysed trimethylsilylazide may explosively decompose.

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A process for an azide-free preparation of imine ligands (i.e. phosphinimine) is described in Canadian patent application CA 2,261,518. However, this procedure encompasses two reaction steps starting from aminophosphoniumhalides. Another disadvantage of the method described in CA 2,261,518, is the use of harmful and costly reagents, such as *n*-butyllithium. Finally, in this procedure the imine ligand is substituted with trimethylsilylchloride, which is removed as such in a subsequent reaction of the imine ligand with the metal-organic precursor. Known production processes for guanidine-, ketimine- and iminoimidazoline comprising metal-organic compounds are described in WO 2070569 and US 6114481. They are carried out at low temperature and require in some cases a solvent change.

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Disadvantage of the known less dangerous method is thus that at least two steps are required, when starting the process with an aminophosphoniumhalide. Purpose of the present invention is to provide a widely applicable method for the manufacturing of a metal-organic compound from an imine and a metal-organic precursor in one step.

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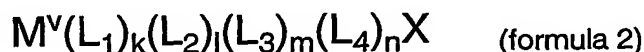
This aim is achieved in that an imine ligand according to formula 1, or the HA adduct thereof, wherein HA represents an acid, of which H represents its proton and A its conjugate base, is contacted with a metal-organic reagent of formula 2 in the presence of at least 1, respectively 2 equivalents of base, wherein

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wherein Y is selected from a substituted carbon, nitrogen or phosphorous atom and R represents a protic or aprotic substituent, and:

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wherein:

M represents a group 4 or group 5 metal ion

15 V represents the valency of the metal ion, being 3, 4 or 5

L_1 , L_2 , L_3 , and L_4 represent ligands on M and may be equal or different

X represents a group 17 halogen atom

$k, l, m, n = 0, 1, 2, 3, 4$ with $k+l+m+n+1=V$

20 With the method of the invention a metal-organic compound, suitable as precatalyst in olefin polymerisation, is prepared in one step. An additional advantage of the method of the invention is, that during the process hardly any by-products are formed, so that further purification is not necessary (or very limited with respect to state of the art processes). The metal-organic compound prepared by the method of the invention has a higher purity than a metal-organic compound prepared via known
25 production processes and can be used as such in olefin polymerisation processes. An additional advantage of the process of the invention is that the process can be carried out at room temperature, whereas the reaction of the N-trialkylsilyl substituted imine ligand with the metal-organic reagent has to be often carried out at elevated temperatures.

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The imine derivative or its HA adduct, as represented in formula 1, is substituted by an Y-and an R group. In the method of the invention, the Y group consists of a substituted carbon, nitrogen or phosphorous atom. If Y represents a substituted carbon atom, the number of substituents is 2. If Y represents a substituted nitrogen atom, the number of substituents is 1 and the number of substituents is 1 or 3 if

Y represents a phosphorous atom, depending on the valency of the phosphorous atom. Substituents on carbon, nitrogen or phosphorous may be equal or different, optionally linked with each other, optionally having heteroatoms. Substituents may be protic or aprotic. A protic substituent is defined here as a substituent, which has at least one,
 5 group 15 or group 16 atom containing at least one proton.

Examples of protic substituents include C₁-C₂₀ linear, branched or cyclic hydrocarbyl radicals, substituted with a group 15 or 16 atom bearing at least one hydrogen atom. Preferred protic substituents include phenolic radicals, pyrrolic radicals, indolic radicals, and imidazolic radicals.

10 The substituent is called aprotic if the substituent lacks a group containing a group 15 or group 16 atom bearing a proton. An unsubstituted aprotic hydrocarbyl radical can be a C₁-C₂₀ linear, branched or cyclic radical, a hydrogen atom, a halogen atom, a C₁₋₈ alkoxy radical, a C₆₋₁₀ aryl or aryloxy radical, an amido radical, or a C₁₋₂₀ hydrocarbyl radical unsubstituted or substituted by a halogen atom, a
 15 C₁₋₈ alkoxy radical, a C₆₋₁₀ aryl or aryloxy radical, an amido radical, a silyl radical of the formula 4, or a germanyl radical of the formula 5.

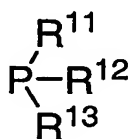
The substituent R can be H, or being equal as these for the substituent on Y.

Examples of imine ligands according to formula (1) thus include:
 20 guanidines, iminoimidazolines, phosphinimines, phenolimines, pyrroleimines, indoleimines and imidazoleimines.

R may be linked with Y, thus forming a ring system, optionally comprising heteroatoms, or optionally comprising functional groups. Examples of ligands comprising such ring systems include: 8-hydroxyquinoline, 8-aminoquinoline, 8-
 25 phosphinoquinoline, 8-thioquinoline, 8-hydroxyquinaldine, 8-aminoquinaldine, 8-phosphinoquinaldine, 8-thioquinaldine and 7-azaindole or indazole.

In a preferred embodiment of the method of the invention, R represents a hydrogen atom and Y is selected from the group consisting of:

30 i) a phosphorus substituent according to the formula:



(formula 3)

wherein each R^{1j} , with $j = 1-3$ is independently selected from the group consisting of a hydrogen atom, a halogen atom, a C_{1-8} alkoxy radical, a C_{6-10} aryl or aryloxy radical, an amido radical, or a C_{1-20} hydrocarbyl radical unsubstituted or substituted by a halogen atom, a C_{1-8} alkoxy radical, a C_{6-10} aryl or aryloxy radical, an amido radical, a silyl

5 radical of the formula:



or a germanyl radical of the formula:

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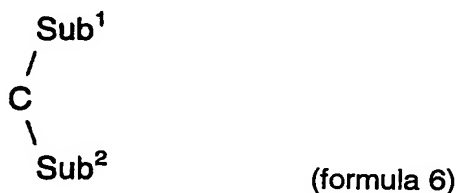


wherein R^{2j} , with $j=1-3$, is independently selected from the group consisting of hydrogen, a C_{1-8} alkyl or alkoxy radical, C_{6-10} aryl or aryloxy radicals,

15 each substituent R^{1j} or R^{2j} may be linked with another R^{1j} or R^{2j} respectively to form a ring system,

ii) a substituent according to formula 6:

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wherein each of Sub^1 and Sub^2 is independently selected from the group consisting of hydrocarbyl radicals having from 1 to 30 carbon atoms; silyl radicals, (substituted) amido radicals and (substituted) phosphido radicals, and wherein Sub^1 and Sub^2 may be linked with each other to form a ring system.

Preferably Sub^1 and Sub^2 are each independently selected from the group of C1-C20

hydrocarbonyl radicals, or substituted amido radicals optionally linked by a bridging moiety.

Preferably the metal-organic compound is contacted with the HA adduct of an imine ligand, with more preference a phosphinimine ligand in the presence of at least two equivalents of a base. A process for a less dangerous azide-free preparation of phosphinimine ligands is described in Canadian patent application CA 2,261,518. In a first step a trialkyl aminophosphoniumhalide (which is the HX adduct of trialkylphosphinimine) is reacted with a base to a trialkyl phosphinimide salt, where after this trialkyl phosphinimide salt is quenched with trimethylsilylchloride(TMSCl). The thus formed trialkyl silyl substituted phosphinimine ligand is subsequently reacted in a third step with CpTiCl_3 to the metal organic compound. A disadvantage of the 3 step method described in CA 2,261,518, is the use of harmful and costly reagents, such as *n*-butyllithium. Another purpose of the present invention is therefore to provide a widely applicable method for the manufacturing of a metal-organic compound from an aminophosphoniumhalide and a metal-organic reagent in one step. This aim is achieved in a process wherein an HA adduct of a phosphinimine ligand (e.g. aminophosphonium halide) according to formula 1 is contacted with a metal-organic reagent of formula 2 in the presence of at least 2 equivalents of a base, wherein HA represents an acid, of which H represents its proton and A its conjugate base, with $\text{Y}=\text{NH}$ being the ligand in formula 1, wherein Y is represented by formula 3.

In the process of the invention, HA represents an acid, of which H represents its proton and A its conjugate base. Examples of A are halogenides, such as fluoride, chloride, bromide, or iodide, sulfate, hydrogensulfate, phosphate, hydrogenphosphate, dihydrogenphosphate, carbonate, hydrogencarbonate, aromatic or aliphatic carboxylates, cyanide, tetrafluoroborate, (substituted) tetraphenylborates, fluorinated tetraarylborates, alkyl or aryl sulfonates.

The "number of equivalents of a base" is understood to be the amount of equivalents with respect to the number of imine ligands, or functionalities in the event that one ligand comprises more than one imine functionality. With "at least 1, respectively 2 equivalents of a base", and later on in the application "at least 3, respectively 4 equivalents of a base", is meant that at least 1, respectively 3 equivalents of a base are required when the imine ligand as such is used, but that at least 2, respectively 4 equivalents are required, in case the HA adduct of the imine ligand is used.

The metal-organic reagent used in the method of the invention is a reagent according to formula 2. In this formula L_1 to L_4 can independently be a monoanionic ligand or a group 17 halogen atom.

Examples of monoanionic ligands are: halides like a fluoride,
5 chloride, bromide or iodide, (un)substituted aliphatic or aromatic hydrocarbyls, like C_1 - C_{20} hydrocarbyl radicals, aryloxy or alkyloxy, cyclopentadienyls, indenyls, tetrahydroindenyls, fluorenyls, tetrahydrofluorenyls, and octahydrofluorenyls, amides, phosphides, sulfides, ketimides, guanidines, iminoimidazolines, phosphinimides, substituted imines, like (hetero)aryloxyimines, pyrroleimines, indoleimines,
10 imidazoleimines or (hetero)aryloxides.

Preferred monoanionic ligands include: fluoride, chloride, bromide, iodide, C_1 - C_{20} hydrocarbyl radicals, cyclopentadienyl, C_1 - C_{20} hydrocarbyl substituted cyclopentadienyls, halogen substituted C_1 - C_{20} hydrocarbyl substituted cyclopentadienyls, indenyl, C_1 - C_{20} hydrocarbyl substituted indenyls, halogen
15 substituted C_1 - C_{20} hydrocarbyl substituted indenyls, fluorenyls, C_1 - C_{20} hydrocarbyl substituted fluorenyls, halogen substituted C_1 - C_{20} hydrocarbyl substituted fluorenyls, C_1 - C_{45} substituted phosphinimides, C_1 - C_{20} substituted ketimides, C_1 - C_{30} substituted guanidines, C_1 - C_{30} iminoimidazolines.

Most preferably monoanionic ligands are selected from fluoride,
20 chloride, bromide, iodide, cyclopentadienyl, C_1 - C_{20} hydrocarbyl (optionally containing hetero- or group 17 halogen atoms), substituted cyclopentadienyls, indenyl, C_1 - C_{20} hydrocarbyl substituted indenyls, and halogen substituted C_1 - C_{20} hydrocarbyl substituted indenyls.

Depending on the valency of the metal of the metal-organic reagent,
25 preferably at least one L_1 , L_2 , L_3 , or L_4 represents a group 17 atom. If the valency of the metal $V = 3$, one or two ligands L may represent a group 17 atom. If $V = 4$, two or three ligands L may represent a group 17 atom. If $V = 5$, two to four ligands L may represent a group 17 atom. Preferred group 17 atom ligands are fluoride, chloride, bromide or iodide atoms. The most preferred group 17 atom ligand is chloride. In a most preferred
30 embodiment, at least one of the ligands L is chosen from cyclopentadienyl, C_1 - C_{20} hydrocarbyl (optionally containing hetero- or group 17 halogen atoms), substituted cyclopentadienyls, indenyl, C_1 - C_{20} hydrocarbyl substituted indenyls, and halogen substituted C_1 - C_{20} hydrocarbyl substituted indenyls. C_1 - C_{20} hydrocarbyl (optionally containing hetero- or group 17 halogen atoms) also includes aryloxy or alkyloxy,
35 octahydrofluorenyls, amides, phosphides, sulfides, ketimides, guanidines,

iminoimidazolines, phosphinimides, substituted imines, like (hetero)aryloxyimines, pyrroleimines, indoleimines, imidazoleimines and (hetero)aryloxides.

In the method of the invention an imine ligand or the HA adduct thereof according to formula 1, is contacted with a metal-organic reagent of formula 2
5 in the presence of at least 1, respectively 2, equivalents of a base. Examples of a base include, carboxylates (for example potassium acetate), fluorides, hydroxides, cyanides, amides and carbonates of Li, Na, K, Rb, Cs, ammonium and the group 2 metals Mg, Ca, & Ba, the alkali metal (Li, Na, K, Rb, Cs) phosphates and the phosphate esters (eg. $C_6H_5OP(O)(ONa)_2$ and related aryl and alkyl compounds) and their alkoxides and
10 phenoxides, thallium hydroxide, alkylammonium hydroxides and fluorides. Some of these bases may be used in conjunction with a phase transfer reagent, such as for example tetraalkylammonium-, tetraalkylphosphonium salts or crown ethers. Also stronger bases may be applied, like carbanions such as hydrocarbanions of group 1, group 2, group 12 or group 13 elements. Also the metallic alkalimetals of group 1
15 may be applied as a base.

Preferred bases include amines, phosphanes, organolithium compounds, or organomagnesium compounds, alkali metals, group 1 hydrides or group 2 hydrides

More preferred bases are mono-, di-, or tri-, alkylamines or aromatic
20 amines, organolithium compounds, organomagnesium compound, sodium hydride or calciumhydride. Under aromatic amines is understood in this application compounds having a nitrogen atom in an aromatic ring system or mono-, di-, or triarylamines.

Even more preferred bases are triethylamine, pyridine, tripropylamine, tributylamine, 1,4-diaza-bicyclo[2.2.2]octane, pyrrolidine or piperidine
25 organolithium compounds, or organomagnesium compounds. Examples of organomagnesium compounds are: methylmagnesiumhalides, phenylmagnesiumhalides, benzylmagnesiumhalides, biphenylmagnesiumhalides, naphthylmagnesiumhalides, tolylmagnesiumhalides, xylylmagnesiumhalides, mesitylmagnesiumhalides, dimethylresorcinolmagnesiumhalides, N,N-
30 dimethylanilinemagnesiumhalides, dimethylmagnesium, diphenylmagnesium, dibenzylmagnesium, bis(biphenyl)magnesium, dinaphthylmagnesium, ditolylmagnesium, dixylylmagnesium, dimesitylmagnesium, bis(dimethylresorcinol)magnesium, bis(N,N-dimethylaniline)magnesium.

Examples of organolithium compounds are: methyllithium,
35 phenyllithium, benzylolithium, biphenyllithium, naphthyllithium, dimethylresorcinollithium,

N,N-dimethylanilinelithium.

In order to make a polyolefin by a borane or borate activatable metal-organic compound, the halide groups of the metal-organic compound from the process of the invention have to be alkylated or arylated in an additional reaction step. This can
5 be done for example with an organolithium compound or an organo magnesium compound. Surprisingly it has been found that such alkylated or arylated metal-organic compound can also be prepared in one step by the process of the invention by carrying out the process in the presence of at least 3, respectively 4 equivalents of an organomagnesium compound or an organolithium compound as a base. This holds for
10 a metal-organic reagent comprising 3 halogen ligands reacting with 1 imine functionality only. One skilled in the art will understand that metal-organic reagents with 4 or 5 halogen ligands will require at least 4 respectively 5 equivalents of a base in stead of at least 3; or 5 respectively 6 equivalents in stead of 4.

The process of the invention is preferably carried out in a solvent.
15 Suitable solvents are solvents that do not react with the metal-organic reagent or the metal-organic compound formed in the process of the invention. Examples of suitable solvents include aromatic and aliphatic hydrocarbons, halogenated hydrocarbons, amides of the aliphatic carboxylic acids and primary, or secondary amines, DMSO, nitromethane, acetone, acetonitrile, benzonitrile, ethers, polyethers, cyclic ethers, lower
20 aromatic and aliphatic ethers, esters, pyridine, alkylpyridines, cyclic and primary, secondary or tertiary amines, and mixtures thereof. Preferred solvents include aromatic or aliphatic hydrocarbons or mixtures thereof.

The process of the invention can be carried out, by adding at least 1, respectively at least 2 equivalents of a base to a mixture of the imine ligand or its HA
25 adduct and the metal-organic reagent thus forming a reaction mixture. The desired metal-organic compound is often formed instantaneously. Excess of a base may be applied without negative effects on the reaction product. However in the event that the base is an organic base, the process of the invention is preferably carried out in the presence of 1 equivalent of an organic base with respect to the imine ligand. An
30 advantage of a process of the invention in the presence of only one equivalent of a base is that the resulting compound may have a higher activity. Without being bound to an explanation, this may be a consequence of the fact that the formation of a coordination complex of the organic base with the metal-organic compound is prevented. An organic base is understood to be a neutral organic compound, capable
35 of accepting a proton. In this application an inorganic base is understood to be a metal

or a metal salt capable of accepting at least one proton. A metal-organic base is a carbanion of an element of group 1,2,12 or 13. Preferably the application is carried out in the presence of an inorganic base or a metal-organic base. An advantage of the use an inorganic base is the formation of a precipitate which can be filtered off easier than
5 the precipitate which is formed with the organic base used in CA 02243775. Still another advantage is the fact that inorganic bases are environmentally friendly and allows a broader range of solvents used in the process of the invention. An advantage of the metal-organic base is that with this base the hydrocarbylated metal-organic compound can be prepared in a one step process. In the known process the metal-
10 organic dichloride has to be hydrocarbylated in a second reaction step in case this is required for example to use the metal-organic compound in a polymerisation process using a boron comprising activator. Surprisingly it turned out that the reaction even with only one equivalent of an organic base appeared to be instantaneously at room temperature to quantitative conversion. Another advantage of a process of the
15 invention in the presence of only one equivalent of a base, is that the resulting compound may have a higher activity. Without being bound to an explanation, this may be a consequence of the fact that the formation of a coordination complex of the organic base with the metal-organic compound is prevented.

During the reaction, a salt is formed. The reaction mixture as
20 obtained by contacting an imine or its HA adduct may be used as precatalyst in a polyolefin polymerisation without an additional purification step if the salt formed during the reaction is compatible with the polymerisation process. A purification step can be a filtration, a crystallisation or a precipitation. Preferably the purification is a filtration step. If a salt free metal-organic compound is required, the salt can be removed by using a
25 filtration. Depending on the solubility of the metal-organic compound, the mixture may be heated and then filtered. An advantage of the present invention is that the filtrate may be used as such without further purification in a following process, such as an alkylation or arylation step or the polymerisation process. If desired, the metal-organic compound may be isolated by distillation of the solvent, by precipitation or by
30 crystallisation from a suitable solvent.

The invention further relates to a process for the preparation of a polyolefin in the presence of an activator wherein the process is carried out in the presence of a metal-organic compound comprising at least one imine ligand according to formula 1 of claim 1, obtained by a process wherein an imine, or the HA adduct
35 thereof, wherein HA represents an acid, of which H represents its proton and A its

conjugate base, is contacted with a metal-organic reagent of formula 2 of claim 1 in the presence of at least 1 equivalent, respectively at least 2 equivalents of a base.

Preferably R represents a hydrogen atom and wherein Y is selected from the group consisting of: a phosphorus substituent according to formula 3 or a substituent

5 according to formula 6. With more preference the process is carried out in the presence of at least 3, respectively at least 4 equivalents of an organomagnesium- or an organolithium compound, as the base. Such an olefin polymerisation can be carried out in solution, slurry or in the gas phase. An activator is understood to be a co-catalyst as described in *Chem. Rev.*, **2000**, *100*, 1391 by E. Y-X. Chen and T.J. Marks.

10 In a preferred embodiment of the olefin polymerisation the (alkylated) metal-organic compound is formed in situ. By in situ preparation is meant in this context, that the metal-organic compound is made and subsequently activated in or anywhere before the reactor of the polymerisation equipment by contacting an imine or its HA adduct with an metal-organic reagent in the presence of an olefin polymerisation
15 compatible base.

In the in situ preparation of the metal-organic compound, it turned out to be favourable to use a surplus of ligand. The number of ligands which are effectively bound to the metal ion are then determined by the number of equivalents of the base. In this case the "number of equivalents of a base" should then be read as the number
20 of equivalents of a base with respect to the equivalents of the ligands being bound to the metal ion. Examples of bases compatible with the olefin polymerisation process include amines, organomagnesium compounds, organolithium reagents, organozinc reagents, organoaluminum reagents. More preferred bases are: aromatic amines, organomagnesium compound, organolithium reagents, organozinc reagents,
25 organoaluminum reagents. Most preferred bases are N,N-dimethylaniline, diphenylmethylamine, triphenylamine, dibutylmagnesium, n-butyllithium, C₁-C₂₀ dihydrocarbylzinc derivatives, diisobutylaluminium hydride, C₁-C₂₀ trihydrocarbyl aluminiums, or aluminoxanes. In the case where aluminoxanes are applied as a base, the base can be the activator.

30 In the olefin polymerisation according to the invention, R preferably represents a proton and Y is preferably selected from the group consisting of:

- i) a phosphorus substituent according to formula 3 of claim 2 or,;
- ii) a substituent according to formula 6.

Advantages of the process of the invention are: mild conditions,
35 higher yields, higher reaction rates and smaller amounts of by-products. The (alkylated)

metal-organic compounds as obtained by the invented process can be used without further purification in the olefin polymerisation resulting in more active catalysts.

The invention will be elucidated with some non-limiting examples:

5 General part

Experiments were performed under a dry and oxygen-free nitrogen atmosphere using Schlenk-line techniques. ^1H -NMR, ^{13}C -NMR-spectra and ^{31}P -NMR-spectra were measured on a Bruker Avance 300 spectrometer. Diethyl ether and ligroin were distilled from sodium/potassium alloy; THF and toluene from potassium and sodium, respectively, all having benzophenone as indicator.

10 Tri-ethylamine was distilled from calciumhidoide before use.

Other starting materials were used as obtained.

Comparative experiment ASynthesis of N-trimethylsilyltri-*tert*-butylphoshinimine

To neat tri-*tert*-butylphosphane (4.38 g, 21.7 mmol) was added azidotrimethylsilane (1.00 mL, 0.87 g, 7.56 mmol). The mixture was heated to the temperature where the formation of nitrogen started (approximately 110°C). A white precipitate started to form. The remaining amount of azide (1.62 g, 14.1 mmol) was added portionwise in order to control the reaction. The product was distilled resulting in 4.20 g (66%) of N-trimethylsilyltri-*tert*-butylphoshinimine.

10 Preparation of (Cp)Ti(NP(*t*-Bu)₃)Cl₂ using CpTiCl₃ and N-trimethylsilyltri-*tert*-butylphoshinimine

To a solution of cyclopentadienyltitanium trichloride, CpTiCl₃ (0.430g, 1.96 mmol) in toluene (25 mL) was added solid N-trimethylsilyltri-*tert*-butylphoshinimine (0.566g, 1.96 mmol). The solution was heated to 60°C for 30 minutes and subsequently stirred overnight. The volatiles were removed in vacuo and the product was washed three times with ligroin. Drying of the yellow solid resulted in 0.63 g (81%). Overall yield with respect to the phosphine: 53%.

20 Part A Examples related to the preparation of the preparation of the metal-organic compound.

Example I. One-step preparation of (Cp)Ti(NP(*t*-Bu)₃)Cl₂ from Tri-*tert*-butyl Aminophosphonium Chloride (*t*-Bu₃PClNH₂) and CpTiCl₃ using triethylamine as base

a. Synthesis of t-Bu₃PClNH₂ at atmospheric pressure

25 To a solution of *tert*-butylphosphane (4.06 g, 20.1 mmol) in ether (60 mL) was added hexachloroethane (4.76 g, 20.1 mmol). The mixture became heterogeneous. Acetonitrile (20 mL) was added to obtain a homogeneous solution. ³¹P-NMR showed the oxidation to be complete. Ammonia gas was bubbled through at atmospheric pressure for 20 minutes. After 16 hours, the conversion appeared to be 71% according to ³¹P-NMR. NH₃ was bubbled through again for 10 minutes. The reaction was complete after stirring for 3 days at room temperature and atmospheric pressure. The solvents were removed in vacuo resulting in 4.98 g (98%) of a white powder being characterized by ¹H-NMR and ³¹P-NMR as tris(*tert*)butylaminophosphoniumchloride.

35 *b. One-step preparation of (Cp)Ti(NP(*t*-Bu)₃)Cl₂ from Tri-*tert*-butyl*

Aminophosphonium Chloride (t-Bu₃PClNH₂) and CpTiCl₃ using triethylamine as base.

To a solution of commercially available CpTiCl₃ (0.55 g, 2.5 mmol) in toluene (20 mL) was added the aminophosphoniumchloride prepared under a (0.63 g, 2.5 mmol). To the almost clear solution was added an excess of triethylamine (2.5 mL, 18 mmol). The reaction mixture became more heterogeneous and the colour changed to orange. After stirring the reaction mixture overnight, the formed triethylammoniumchloride was filtered. The solvents from the filtrate were removed in vacuo. NMR analysis (¹H, ³¹P and ¹³C) showed (Cp)Ti(NP(t-Bu)₃)Cl₂ with no detectable amounts of by-product.

Example II. One-step preparation of (Cp-C₆F₅)Ti(NP(t-Bu)₃)Cl₂ from Tri-tert-butyl Aminophosphonium Chloride (t-Bu₃PClNH₂) and Cp(C₆F₅)TiCl₃ using triethylamine as base

To a solution of C₆F₅CpTiCl₃ (1.00 g, 2.59 mmol) (obtained by the method described in *J. Organomet. Chem.*, 2000, 107 by Rausch et. al.) in toluene (60 mL) was added t-Bu₃PClNH₂ (0.68 g, 2.59 mmol). To the orange mixture was added triethylamine (1.0 mL, 7.2 mmol). A precipitate was formed immediately and NMR monitoring of the reaction mixture showed complete conversion to the desired products, with no detectable amounts of by-product. The mixture was stirred for 3 days. The reaction mixture was filtered and the solvent and excess triethylamine were removed in vacuo resulting in 1.22 g (83%) (Cp-C₆F₅)Ti(NP(t-Bu)₃)Cl₂. ³¹P- and ¹H-NMR showed (Cp-C₆F₅)Ti(NP(t-Bu)₃)Cl₂ with no detectable amounts of by-product.

Example III. One-step preparation of (Cp-C₆F₅)Ti(NP(t-Bu)₃)Me₂ from Tri-tert-butyl Aminophosphonium Chloride (tBu₃PClNH₂) and Cp(C₆F₅)TiCl₃ using methylmagnesiumbromide as base

To an orange mixture of C₆F₅CpTiCl₃ (1.00 g, 2.59 mmol) and t-Bu₃PClNH₂ (0.68 g, 2.59 mmol) in toluene (60 mL) and THF (20mL) was added a MeMgBr solution in ether (3.0M, 4.0 mL, 12 mmol) at -20°C. The reaction mixture was stirred for 45 minutes and subsequently dried in vacuo. The residue was extracted with boiling ligroin (20 and 40 mL respectively). The solvents were removed in vacuo resulting in 1.33 g (98%) of (Cp-C₆F₅)Ti(NP(t-Bu)₃)Me₂ with no detectable amounts of by-product.

Example IV. One-step preparation of $\text{CpTi}(\text{NP}(\text{n-Bu})_3)\text{Cl}_2$ from Tri-*n*-butyl Aminophosphonium Bromide ($\text{n-Bu}_3\text{PBrNH}_2$) and CpTiCl_3 using triethylamine as base

a. *Synthesis of $\text{n-Bu}_3\text{PBrNH}_2$*

Tri-*n*-butylphosphane (20.2 g, 0.10 mol) was dissolved in acetonitrile (200mL). The solution was cooled to 0°C and bromine (16.2 g, 0.10 mol) was added in 10 minutes. An exothermic effect was observed. After 10 minutes, the cooling bath was removed. The bright yellow mixture was stirred for 2 hours after reaching room temperature. The mixture was again cooled to 0°C and ammonia was introduced. An exothermal reaction occurred. The temperature increase was controlled by the addition rate of the ammonia. The yellow slurry turns white after 15 minutes and ammonia was bubbled through for an additional 10 minutes. The acetonitrile was removed in vacuo and the residue was extracted with dichloromethane (2x150ml). The solution was decanted from the solids and the solvent was subsequently removed in vacuo resulting in a white solid. Yield: 28.2 g (95%) $\text{n-Bu}_3\text{PBrNH}_2$.

b. *One-step preparation of $\text{CpTi}(\text{NP}(\text{n-Bu})_3)\text{Cl}_2$ from Tri-*n*-butyl Aminophosphonium Bromide ($\text{n-Bu}_3\text{PBrNH}_2$) and CpTiCl_3 using triethylamine as base*

CpTiCl_3 (2.21g, 10.1 mmols) and $\text{n-Bu}_3\text{PNH}_2\text{Br}$ (3.05g 10.2 mmols), were dissolved in toluene (80mL). At room temperature triethylamine (4mL, 29 mmol) was added dropwise over a period of 10 minutes. The reaction mixture immediately became heterogeneous and the colour changed from orange to bright yellow. The mixture was stirred for 1 hour at room temperature (according to ^{31}P NMR the reaction was converted completely to the desired product). The ammoniumsalt was filtered off and washed once with 25 mL of toluene. The solvent was subsequently removed in vacuo leaving a viscous residue (product contaminated with small amounts of solvent). In order to obtain a solid product, the resulting residue was dissolved in 80 ml of hexanes and 25 mL of dichloromethane. Removing the solvent in vacuo yielded 3.4g of the product as a yellow solid (85%) being $\text{CpTi}(\text{NP}(\text{n-Bu})_3)\text{Cl}_2$.

Example V. Synthesis of tris(*N,N*-dimethylamido)phosphoraneimido cyclopentadienyl titanium(IV) dichloride

To a cold solution (−60°C) of cyclopentadienyltitanium trichloride (0.50g, 2.28 mmol) in toluene (30 mL) was added

N,N,N',N',N'',N''-hexamethylphosphorimidic triamide (0.41 g, 2.3 mmol). The mixture was allowed to warm to room temperature. Then, triethylamine (1.0 mL, 7.2 mmol) was added. A precipitate formed directly after the addition of the triethylamine. ³¹P-NMR reaction monitoring showed that the desired product was formed without any
5 detectable amount of by-product. The reaction mixture was filtered in order to remove the triethylammonium chloride. The solvents were removed in vacuo and the residue was crystallised from toluene to give 0.73 g (yield: 89%) of a yellow crystalline product, which was characterized by ¹H- and ³¹P-NMR to be tris(N,N-
dimethylamido)phosphoraneimido cyclopentadienyl titanium(IV) dichloride.

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Example VI. Synthesis of 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline
cyclopentadienyl titanium dichloride

To a suspension of 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline
(1.50 g, 5.0 mmol) (prepared according to the procedure by L.Toldy et al, US Patent
15 4,284,642), and cyclopentadienyltitanium trichloride (1.10 g, 5 mmol) in toluene (80 mL) was added triethylamine (1.0 mL, 7.2 mmol) at ambient temperature. After stirring for 1 hour, the suspension was heated to reflux, then filtered hot. Cooling to ambient temperature gave orange crystals, which were filtered, washed with cold toluene and dried (1.36 g, 57% yield). Partial evaporation of the mother liquor and cooling to -20 °C
20 afforded another 0.90 g (38%). Total yield of 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline cyclopentadienyl titanium dichloride was 95%.

Example VII. Synthesis of 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline
cyclopentadienyl titanium dimethyl

To a suspension of 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline
(5.86 g, 20.0 mmol) and cyclopentadienyltitanium trichloride (4.39 g, 20.0 mmol) in toluene (200 mL) was added triethylamine (2.53 g, 25 mmol) at ambient temperature. After stirring for 1 hour at ambient temperature, the thick yellow-orange suspension was heated to reflux and filtered hot. The yellow residue was extracted with boiling
30 toluene portions of 10 mL 4 times (leaving a grey-white residue). The combined orange filtrates (separating yellow-orange crystals upon cooling) were cooled to 0°C. Methyl magnesium bromide (14 mL of a 3.0 M solution in diethyl ether, 44 mmol) was added in 10 minutes. The orange suspension turned yellow gradually. The mixture was stirred overnight, then evaporated to dryness. The residue was extracted with boiling ligroin
35 (200 mL) and the resulting suspension was filtered hot. Cooling to approx. -20°C

afforded yellow crystals, which were filtered and washed with cold ligroin to give 2.8 g (32% yield) of NMR pure product. From the partially evaporated mother liquor and 2nd ligroin extract, a 2nd fraction of pure product was obtained (1.0 g, 11%). Total yield of 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline cyclopentadienyl titanium dimethyl was 43%.

Example VIII. Synthesis of 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline cyclopentadienyl titanium dimethyl using methylmagnesium bromide as base.

To a suspension of 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline (2.93 g, 10.0 mmol) and cyclopentadienyltitanium trichloride (2.19 g, 10.0 mmol) in toluene (100 mL) was added methylmagnesiumbromide (11 mL of a 3.0 M solution in diethyl ether, 33 mmol) at -80°C during 10 minutes. The mixture was allowed to warm to ambient temperature to give a yellow suspension. THF (30 mL) was added, and the mixture was stirred for 15 hours. The light yellow suspension was evaporated to dryness. The residue was extracted with boiling ligroin (100 mL). The resulting suspension was filtered hot. The cake was extracted further with hot ligroin (Three times with 60 mL until the filtrate became colourless). The combined yellow filtrates were partially evaporated under reduced pressure to 50 mL. Cooling to approx. 4°C afforded yellow crystals, which were filtered and washed with cold ligroin to give 2.05 g (47% yield) of NMR pure 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline cyclopentadienyl titanium dimethyl.

Example IX. Synthesis of 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline cyclopentadienyl titanium dichloride

a. Synthesis of 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline

To a mixture of 2,6-diisopropylaniline (260 g, 1.47 mol) in ethanol (1200 mL) was slowly added glyoxal (108.5 g of a 40 w-% in water solution, 0.75 mol). The solution turned intensely red, then intensely yellow. The mixture was heated to reflux overnight. Cooling to 4 degrees resulted in crystallisation of yellow material, which was filtered and washed with cold ethanol until filtrate became bright yellow (instead of brown). The bright yellow powder was dried (202.6 g, 72%). This diimine (100 g, 0.27 mol) was dissolved in ethanol (1000 mL). The mixture was cooled to 0°C. Sodium borohydride (102.1 g, 2.7 mol) was added in portions during 1 hour. The mixture was allowed to warm to room temperature, then stirred 1 hour. The mixture was heated to reflux gently (gas evolution!) and heated to reflux for 1

hour. After cooling, the mixture was admixed with water (2L), and the suspension filtered. The yellow precipitate was dried (100.1 g, 98%).

57 g (0.15 mol) of the diamine was dissolved in toluene (250 mL) and heated to reflux. A solution of cyanogen bromide (19.1 g, 0.18 mol) in toluene (100 mL) was added during the course of ~1 hour, resulting in formation of a grey precipitate in an orange-red solution. After stirring at reflux for 1 hour, the mixture was cooled. The precipitate was filtered, washed with toluene and ligroin (to give 47.1 g yellow light powder). This powder was dissolved in water/ethanol 400/500 mL, and 10.0 M NaOH in water was added until strongly basic (pH>10). The precipitate was filtered and washed with water, then dried to give 37.3 g (61.4% yield) of near pure product. The iminoimidazoline can be crystallized to give pure material as colourless crystals from boiling ligroin (270 mL) and filtering hot to remove some insoluble material (recovery 67%).

b. *Synthesis of 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline cyclopentadienyl titanium dichloride*

To a suspension of 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline (1.02 g, 2.5 mmol) and cyclopentadienyltitanium trichloride (0.55 g, 2.5 mmol) in toluene (20 mL) was added triethylamine (0.4 mL, 4.0 mmol) at ambient temperature. After stirring for 2 hours, the thick yellow-orange suspension was filtered, and the filtrate evaporated to dryness to afford 1.31 g (89% yield) of NMR-pure 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline cyclopentadienyltitanium dichloride.

c. *Synthesis of 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline cyclopentadienyl titanium dichloride (reversed addition)*

The same result as under b. was obtained when cyclopentadienyltitanium trichloride and triethylamine were admixed in toluene, and then ligand was added.

d. *Synthesis of 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline cyclopentadienyl titanium dichloride*

To a suspension of 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline (2.03 g, 5.0 mmol) and cyclopentadienyltitanium trichloride (1.10 g, 5.0 mmol) in toluene (30 mL) was added triethylamine (0.8 mL, 5.7 mmol) at ambient temperature. After stirring for 1 hour, the thick yellow-orange suspension was diluted with toluene (50 mL) and ligroin (120 mL). The suspension was heated to reflux and filtered hot. Cooling to approx. 4°C afforded yellow crystals, which were filtered and washed with cold ligroin to give 1.34 g (46% yield) of NMR pure 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline cyclopentadienyl titanium dichloride.

Example X. Synthesis of 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline
cyclopentadienyl titanium dimethyl

To a suspension of 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline (2.06 g, 5.0 mmol) and cyclopentadienyltitanium trichloride (1.10 g, 5.0 mmol) in
5 toluene (40 mL) was added triethylamine (0.8 mL, 5.7 mmol) at ambient temperature. After stirring for 2 hours, the thick yellow-orange suspension was filtered, and the residue washed with toluene. The clear and orange filtrate was partially evaporated (~10 mL solvent removed). After cooling to -78 °C (dry ice/acetone), methyl
10 magnesium bromide solution (3.3 mL of a 3M solution in diethyl ether, 10.0 mmol) was added. The temperature of the mixture was allowed to rise to ambient temperature and the mixture was stirred overnight. The yellow suspension was evaporated to dryness. The residue was extracted with boiling ligroin (80 mL) and the resulting suspension was filtered hot. Evaporation to ~30 mL and cooling to approx. 4°C afforded yellow
15 crystals, which were filtered and washed with cold ligroin to give 1.38 g (51% yield) of NMR pure product. From the partially evaporated mother liquor, a 2nd fraction of pure 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline cyclopentadienyl titanium dimethyl was obtained (0.58 g, 19%). Total yield of 1,3-bis(2,6-diisopropylphenyl)-iminoimidazoline cyclopentadienyl titanium dimethyl: 70%.

20 Example XI. Synthesis of tris(N,N-dimethylamido)phosphoraneimido cyclopentadienyl
titanium(IV) dichloride using potassium carbonate.

To a solution of cyclopentadienyltitanium trichloride (0.51 g, 2.3 mmol) in toluene (40 mL) was added N,N,N',N',N'',N''-hexamethylphosphorimidic triamide (0.41 g, 2.3 mmol). Then, dry K₂CO₃ (0.5 g, 3.6 mmol) was added. ³¹P-NMR
25 reaction monitoring showed that the desired product was formed without any detectable amount of by-product. The reaction mixture was filtered in order to remove the salts which were subsequently extracted with an extra portion of toluene (25 mL). The combined solvents of the filtrate were removed in vacuo to give 0.79 g (yield: 94%) of a yellow crystalline product, which was characterized by ³¹P-NMR to be pure
30 tris(N,N-dimethylamido)phosphoraneimido cyclopentadienyl titanium(IV) dichloride. Surprisingly the process of the invention can be carried out with an inorganic base like potassium carbonate despite of the fact that it has been expected that water may be formed.

35 Example XII. Synthesis of tris(N,N-dimethylamido)phosphoraneimido cyclopentadienyl

titanium(IV) dimethyl

To a solution of cyclopentadienyltitanium trichloride (0.51 g, 2.3 mmol) and N,N,N',N',N'',N''-hexamethylphosphorimidic triamide (0.44 g, 2.5 mmol) in toluene (40 mL) and THF (10 mL) was added a solution of methylmagnesium bromide in ether (2.3 mL, 3.0 M, 6.9 mmol) at room temperature. The reaction was exothermal under gas evolution and the colour changed to light yellow. ³¹P-NMR reaction monitoring showed that the desired product was formed without any detectable amount of by-product. The solvents were removed in vacuo and the product was extracted from the residue with *n*-hexane twice (50 mL each). The solvents were removed in vacuo to give 0.59 g (yield: 79%) of a yellow powder, which was characterized by ¹H- and ³¹P-NMR to be tris(N,N-dimethylamido)phosphoraneimido cyclopentadienyl titanium(IV) dimethyl.

Example XIII. Synthesis of Bis(1-N-cyclohexylcarboximino-6-t-butylphenoxy)titaniumdichloride

To a solution of titanium(IV)chloride (5mL, 1.0M in toluene, 5.0 mmol) in toluene (40 mL) was added 1-N-cyclohexylcarboximine-6-t-butylphenol (2.59g, 10.0 mmol) and triethylamine (1.02g, 10 mmol) subsequently. The reaction mixture was stirred for 16 hours at room temperature. The solid was allowed to precipitate and the supernatant was decanted. The remaining solid was extracted twice with a mixture of toluene/THF (80 mL, 1/1, V/V). The solvents were removed in vacuo resulting in 2.80g (88%) of a red solid. NMR data were consistent with those reported in EP0874005, but the yield of 88 % is substantially higher than the 18 % yield reported in EP 0874005.

Example XIV. Synthesis of Bis(1-N-cyclohexylcarboximino-6-t-butylphenoxy)zirconiumdichloride

To zirconium(IV)chloride (1.40g, 4.5 mmol) was added THF(40mL). The mixture was cooled to 0°C and a solution of 1-N-cyclohexylcarboximine-6-t-butylphenol (2.31g, 8.9 mmol) in toluene (25 mL) was added. Then, triethylamine (0.89 g, 8.9 mmol) was added and the mixture was stirred for 15 hours at room temperature. The solids were allowed to precipitate and the supernatant was decanted from the solid. The solid was extracted with a mixture of toluene/THF (80 mL, 1/1, V/V). The combined extracts were evaporated to dryness resulting in 2.95g (97%) of a light yellow powder. NMR data were consistent with those reported in EP0874005, but the yield was significantly higher than the yield of 43 % reported in EP 0874005.

Part B Examples related to the polymerisation of an olefinic copolymer.

Polymerisation equipment.

The batch copolymerisation was carried out in a polymerisation
5 equipment, having a catalyst dosing vessel equipped with a catalyst dosing pump for
the addition of the catalyst to a 2-liter batch autoclave equipped with a double intermig
stirrer and baffles. The reactor temperature was controlled by a Lauda Thermostat. The
feed streams (solvents and monomers) were purified by contacting them with various
absorption media as is known in the art. During polymerisation, the ethylene (C2) and
10 propylene (C3) were continuously fed to the gas cap of the reactor. The pressure of the
reactor was kept constant by means of a back-pressure valve.

Copolymerisation experiments.

In an inert atmosphere of nitrogen, the reactor was filled with
15 pentamethylheptanes (PMH) (950 mL) and an amount of MAO (Crompton 10wt% in
toluene) and 4-methyl-2,6-di-*tert*-butylphenol (BHT) as given in Tables 1 and 2. The
reactor was heated to 90 °C, while stirring at 1350 rpm. The reactor then was
pressurized to 0.7 MPa and kept under a determined flow of 200 NL/h of ethylene and
400 NL/h of propylene for 15 minutes. Then, the catalyst components were added to
20 the reactor and possible residual material was rinsed with PMH (50 mL) and
subsequently fed to the reactor.

When tritylium tetrakis(perfluorophenyl)borate (TBF20) was used, the
TBF20 was added directly after the catalyst addition. After 10 minutes of
polymerisation, the monomer flow was stopped and the solution was slowly poured into
25 a 2 L Erlenmeyer flask, and dried over-night at 100 °C under reduced pressure.

The polymers were analysed by FT-IR to determine the amount of
incorporated C3 and Intrinsic Viscosity being an indication for the average molecular
weight.

30 Polymer analysis.

The amount of incorporated C3 in weight per cents relative to the
total composition, was measured by means of Fourier transformation infrared
spectroscopy (FT-IR) according to ASTM D 3900 method A.

The Intrinsic Viscosity (IV) was measured at 135°C in decaline.

Examples 1-15: in situ polymerisation

These catalysts were prepared in the polymerisation equipment by adding amounts as depicted in table 1a of toluene solutions of the metal-organic reagent, the ligand and the base successively to the catalyst dosing vessel in toluene
5 (15 mL). After stirring for 5 minutes, the mixture was injected into the polymerisation reactor. Results are shown in Table 1b.

The experiments 1, 2, 5, 12 and 13 were carried out by adding a prepared and purified metal-organic compound to the catalyst dosing vessel, and subsequently fed to the polymerisation reactor.

10 It can be concluded from the comparison of all experiments with experiment 2, that all in situ prepared catalysts produce copolymers having a higher molecular weight than the copolymer produced with the CpTiCl_3 and the base only, which allows preparation of a polyolefin by just adding a metal-organic reagent, an imine ligand and at least 1 equivalent of a base to the polymerisation equipment.

15 From Examples 8 and 10 it can be concluded that a process in the presence of between 5 and 10 equivalents of the imine ligand according to formula 1 is mostly preferred.

Table 1a. In situ polymerisations: polymerisation conditions

Example	Metal-organic reagent/compound	Metal-organic compound dosage ($\mu\text{mol Ti}$)	ligand	Ligand dosage (μmol)	Base	Base dosage (μmol)	Activator system	Al/Ti Molar ratio	BF ₂ O/Ti Molar ratio	BHT/Al Molar ratio	Pol. Time (min)
1	1	0.5	-	-	-	-	MAO/BHT	600		2	10
2	CpTiCl ₃	0.75	-	-	Et ₃ N	0.75	MAO/BHT	3000	-	1	10
3	CpTiCl ₃	1.0	L1	2.0	Et ₃ N	1.0	MAO/BHT	3000	-	1	10
4	CpTiCl ₃	0.4	L1	0.8	Et ₃ N	0.4	MAO/BHT/ TBF ₂ O	3000	2	1	10
5	2	0.05	-	-	-	-	MAO/BHT	3000	-	1	10
6	CpTiCl ₃	0.75	L2	1.5	Et ₃ N	0.75	MAO/BHT	3000	-	1	10
7	CpTiCl ₃	0.75	L2	0.75	Et ₃ N	0.75	MAO/BHT	3000	-	1	10
8	CpTiCl ₃	0.75	L2	3.75	Et ₃ N	0.75	MAO/BHT	3000	-	1	10
10	CpTiCl ₃	0.25	L2	2.5	Et ₃ N	0.25	MAO/BHT	3000	-	1	10
11	CpTiCl ₃	0.4	L2	2	Et ₃ N	0.4	MAO/BHT/ TBF ₂ O	3000	2	1	3
12	TiCl ₄	5	-	-	Et ₃ N	10	MAO/BHT	250	-	1	10
13	3	5	-	-	-	-	MAO/BHT	250	-	1	10

14	TiCl4	5	L3	10	Et3N	10	MAO/BHT	250	-	1	10
15	TiCl4	5	L3	10	-	-	MAO/BHT	250	-	1	10

Metal-organic compound 1 = tris(N,N-dimethylamido)phosphoraneimido cyclopentadienyl titanium(IV) dichloride
Metal-organic compound 2 = 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline cyclopentadienyl titanium dibenzyl
Metal-organic compound 3 = Bis(1-N-cyclohexylcarboximino-6-t-butylphenoxy)titaniumdichloride
L1 =N,N,N',N'',N'''-hexamethylphosphorimidic triamide
L2 =1,3-bis(2,6-dimethylphenyl)-iminoimidazoline
L3 =1-N-cyclohexylcarboximine-6-t-butylphenol

Table 1b. In situ polymerisations: polymerisation results

Example	ΔT (°C)	Yield (g)	residual Ti in polymer (ppm)	Incorpo- rated C3 ⁺ (wt%)	IV (dl/g)
1	0.8	2.93	8.2	41	2.4
2	0.5	2.74	13.1	62	0.96
3	3.5	8.97	5.3	46	Nd
4	1.6	5.34	3.6	42	Nd
5	1.8	6.09	0.4	48	2.77
6	2.0	8.41	4.3	54	2.07
7	0.8	3.76	9.5		
8	4.2	14.37	2.5	51	2.32
10	4.9	19.84	0.6	52	2.29
11	4.4	18.05	1.1	50	
12	0.6	0			
13	1.3	3.55	67.4		1.67
14	1.1	3.18	75.3		1.65
15	1.2	2.89	82.8		1.72

- 5 Examples 17-18: polymerisation with unpurified triisopropylphosphoraneimido cyclopentadienyl titanium(IV) dichloride.

Catalyst preparation

- 10 Cyclopentadienyltitaniumtrichloride (86 mg, 0.39 mmol) and triisopropylaminophosphonium bromide (0.10g, 0.39 mmol) were mixed in toluene (10mL). Triethylamine (80 mg, 0.8 mmol) was added and the reaction mixture was stirred at room temperature for 2 hours.

Polymerisation

For the polymerisation an aliquot (5 mL) of the mixture obtained above, was diluted with toluene (35mL). From this diluted mixture, an aliquot (0.03 mL) was added to the catalyst dosing vessel containing PMH (15 mL). This mixture was subsequently added to the polymerisation reactor and the catalyst dosing vessel was rinsed with PMH (50 mL) .

Examples 19-20: polymerisation with unpurified 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline cyclopentadienyl titanium dibenzyl

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Catalyst preparation

Cyclopentadienyltitaniumtrichloride (75 mg, 0.34 mmol) and 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline (0.10g, 0.34 mmol) were mixed in toluene (10mL). Triethylamine (34 mg, 0.34 mmol) was added and the reaction mixture was stirred at room temperature for 2 hours.

15

Polymerisation

For the polymerisation an aliquot (0.75 mL) of the mixture obtained above was diluted with toluene (25mL). From this diluted mixture, an aliquot (0.15 mL) was added to the catalyst dosing vessel containing PMH (15 mL). This mixture was subsequently added to the polymerisation reactor and the catalyst dosing vessel was rinsed with PMH (50 mL) . Examples 17 – 20 indicate that polymerisation of olefinic monomers is possible by just adding a mixture of a metal-organic reagent, an imine ligand and at least one equivalent of a base to a polymerisation reactor with olefinic monomers, without the need to firstly purify (i.e filtrate) a catalyst (i.e. metal-organic compound) from the mixture.

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Table 2a. Polymerisation with unpurified catalysts: polymerisation conditions

Example	Metal-organic compound	Metal-organic compound dosage ($\mu\text{mol Ti}$)	Activator system	Al/Ti Molar ratio	BF ₂ O/Ti Molar ratio	BHT/Al Molar ratio	Pol. Time (min)
17	4	0.15	MAO/BHT	3000	-	2	10
18	4	0.15	MAO/BHT/TBF ₂ O	3000	2	2	10
19	5	0.15	MAO/BHT	3000	-	1	10
20	5	0.15	MAO/BHT	3000	-	1	10
21	6	0.15	MAO/BHT	300	-	2	10

Metal-organic compound 4 = triisopropylphosphoraneimido cyclopentadienyl titanium(IV) dichloride

- 5 Metal-organic compound 5 = 1,3-bis(2,6-dimethylphenyl)-iminoimidazoline cyclopentadienyl titanium dichloride

Metal-organic compound 6 = triisopropylphosphoraneimido cyclopentadienyl titanium(IV) dimethyl

10

Table 2b. Polymerisation with unpurified catalysts: polymerisation results

Example	ΔT ($^{\circ}\text{C}$)	Yield (g)	residual Ti in polymer (ppm)	Incorporated C ₃ ⁺ (wt%)	IV (dl/g)
17	2.0	6.55	1.1	43	Nd
18	2.7	8.11	0.9	40	Nd
19	4.8	18.75	0.4	501	2.33
20	4.2	16.5	0.4	53	nd
21	3.0	8.82	0.8	43	nd